WHAT IS CLAIMED IS:

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A recombinant adenoviral vector encoding an pro-

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Sub

2. A pharmaceutical composition, comprising the adenoviral vector of claim 1 and a pharmaceutically acceptable carrier.

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3. A method of treating an individual having a pathophysiological state, comprising the step of administering to said individual a pharmacologically effective dose of the composition of claim 2.

4. The method

of claim 3, wherein said

pathophysiological state is a negliartic disease.

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5. The method of claim 4, wherein said neoplastic disease is selected from the group consisting of breast cancer, colon cancer, ovarian cancer, glioma, esteosarcoma and haemopoietic cancers.

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6. The method of claim 4, wherein said composition is administered in a dose of from about 0.1 mg/kg to about 100 mg/kg.

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7. A method of treating an individual having ovarian cancer, comprising the step of administering to said individual a pharmacologically effective dose of a pharmaceutical composition, comprising a recombinant adenoviral vector encoding an proapoptotic bax gene and a pharmaceutically acceptable carrier.

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8. The method of claim 7, wherein said composition is administered in a dose of from about 0.1 mg/kg to about 100 mg/kg.

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chemotherapy and/or radiotherapy in an individual, comprising the step of administering to said individual a pharmacologically effective dose of a pharmaceutical composition, comprising a recombinant adenoviral vector encoding an pro-apoptotic bax gene and a pharmaceutically acceptable carrier.

10. The method of claim 9, wherein said composition is administered in a dose of from about 0.1 mg/kg to about 100 mg/kg.